Data Sheet (Cat.No.T13313)



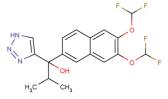
Seviteronel racemate

Chemical Properties

CAS No.: 1375603-36-3 Formula: C18H17F4N3O3

Molecular Weight: 399.34
Appearance: N/A

Storage: 0-4°C for short term (days to weeks), or -20°C for long term (months).



Biological Description

| Description | Seviteronel racemate is the racemate form of Seviteronel and is a potent inhibitor of CYP17 lyase. | |
|----------------------------|--|--|
| Targets(IC ₅₀) | VT-464, h-CYP17 Lyase: 69 nM | |
| In vivo | The MDA-PCa-133 xenograft is derived from a clinical CRPC bone metastasis. Subcutaneous MDA-PCa-133 tumor expresses PSA, full-length androgen receptor (AR), and AR-V7 isoform. The effect of Seviteronel (VT-464) and AA on MDA-PCa-133 growing in tumor-bearing castrated male mice: randomization into three groups; oral treatment with vehicle only, Seviteronel (100 mg/kg bid), or AA (100 mg/kg bid) for 25 days. Both Seviteronel and AA reduced tumor volume (>two fold compared to vehicle; p<0.05). Selective Seviteronel CYP17 lyase inhibition is as effective as AA CYP17 inhibition in this model [2]. | |

Solubility Information

| Solubility | DMSO: 50 mg/mL (125.21 mM) (< 1 mg/ml refers to the product slightly soluble or insoluble) | |
|------------|--|--|
| | | |

Preparing Stock Solutions

| | 1mg | 5mg | 10mg |
|-------|----------|-----------|-----------|
| 1 mM | 2.504 mL | 12.521 mL | 25.041 mL |
| 5 mM | 0.501 mL | 2.504 mL | 5.008 mL |
| 10 mM | 0.25 mL | 1.252 mL | 2.504 mL |
| 50 mM | 0.05 mL | 0.25 mL | 0.501 mL |

Please select the appropriate solvent to prepare the stock solution, according to the solubility of the product in different solvents. The storage conditions and period of the stock solution: - 80 °C for 6 months; - 20 °C for 1 month. Please use it as soon as possible.

Reference

- 1. Rafferty SW, et al. Highly-selective 4-(1,2,3-triazole)-based P450c17a 17,20-lyase inhibitors. Bioorg Med Chem Lett. 2014 Jun 1;24(11):2444-7.
- 2. Sankar N. Maity, et al. Abstract 4772: Efficacy of VT-464, a novel selective inhibitor of cytochrome P450 17,20-lyase, in castrateresistant prostate cancer models. Cancer Research: April 15, 2013; Volume 73, Issue 8, Supplement 1

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